Appl. No. 10/609,143 Amdt. dated February 13, 2006 Reply to Office Action of October 11, 2005

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

1. (currently amended) A method of preparing a composition comprising an agent that affects processing of amyloid precursor protein to beta-amyloid peptide, the method comprising: providing a [[A]] transgenic rodent nonhuman animal or stem cell comprising a diploid genome comprising a transgene encoding a heterologous human APP polypeptide comprising the Swedish mutation wherein the amino acid residues at positions corresponding to positions 595 and 596 in human APP are asparagines and leucine, respectively, wherein the transgene is expressed to produce a human APP polypeptide having the Swedish mutation and wherein the polypeptide is processed to detectable quantities of ATF- β APP in a brain homogenate, which are at least two-fold higher than the quantities of ATF- β APP produced from wild type human β APP in an equivalent transgenic animal;

contacting the transgenic rodent with the agent;

monitoring cleavage of the amyloid precursor protein polypeptide between the Nterminus of the beta amyloid peptide and the ATF-betaAPP in the contacted transgenic rodent
compared to the cleavage in a control transgenic rodent to indicate the agent affects the cleavage;
and

incorporating the agent into a composition with a pharmaceutical carrier.

- 2. (urrently amended) A transgenic nonhuman animal of The method of claim 1, wherein the animal is murine.
 - 3. (canceled)

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- 4. (currently amended) A transgenie nonhuman animal The method of claim 3, wherein the transgene is nonhomologously integrated.
 - 5. (canceled)
- 6. (currently amended) A transgenic nonhuman animal The method of claim 1, wherein the heterologous human APP polypeptide comprising the Swedish mutation is expressed under the transcriptional control of a neural specific enolase promoter.
 - 7-10. (canceled).
- 11. (new) The method of claim 1, wherein the agent inhibits a beta-secretase activity associated with the cleavage.
- 12. (new) The method of claim 1, wherein the dosage of the agent is from 10 μ g/kg to 1 mg/kg.